What is claimed is:

- 1. A method of treating a mammal having type 1 diabetes or at risk for type 1 diabetes, the method comprising administering to the mammal a pharmaceutical composition comprising an agent that inhibits a macrophage migration inhibitory factor (MIF) in the mammal, wherein the agent is a polypeptide or a polynucleotide.
- 2. The method of claim 1, wherein the agent comprises a binding site of an antibody that binds specifically to the MIF.
- 10 3. The method of claim 2, wherein the agent is an antibody.
 - 4. The method of claim 1, wherein the agent is an aptamer that binds specifically to the MIF.
- 5. The method of claim 1, wherein the agent inhibits expression of the MIF.
 - 6. The method of claim 5, wherein the agent is an antisense nucleic acid or mimetic specific for MIF mRNA in the mammal.
- 7. The method of claim 5, wherein the agent is a ribozyme nucleic acid or mimetic specific for MIF mRNA in the mammal.
 - 8. The method of claim 5, wherein the agent is an inhibitory RNA or mimetic specific for MIF mRNA in the mammal.
 - 9. The method of claim 1, wherein the mammal has or is at risk for having diabetes, impaired glucose intolerance, stress hyperglycemia, metabolic syndrome, and/or insulin resistance.
- 30 10. The method of claim 1, wherein the mammal is a rodent.
 - 11. The method of claim 1, wherein the mammal is a human.
- 12. A method of treating a mammal having type 1 diabetes or at risk for type 1
 diabetes, the method comprising administering to the mammal a pharmaceutical composition
 comprising an agent that inhibits a macrophage migration inhibitory factor (MIF) in the
 mammal, wherein the agent is an organic molecule comprising the following structure I or II

$$R_2$$
 R_3
 R_4
 R_4
 R_4
 R_4
 R_4
 R_4

$$R_2$$
 R_3
 R_1
 R_1
 R_1

13. The method of claim 12, wherein the organic molecule comprises structure Π , wherein

X = 0;

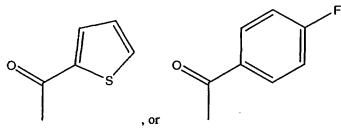
10 Y =

Z = C;

the ring comprising R, and R, =

5

 $R_4 =$



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- 14. A method of evaluating whether a compound is useful for preventing or treating type 1 diabetes, the method comprising
- (a) determining whether the compound inhibits a macrophage migration inhibitory factor (MIF) in a mammal, then, if the compound inhibits the MIF,
 - (b) determining whether the compound inhibits development of type 1 diabetes.
- 15. The method of claim 14, wherein step (b) is performed by evaluating the effect of the compound on proliferation of splenic lymphocytes in the mammal.
- 20 16. The method of claim 14, wherein the compound is a protein.
 - 17. The method of claim 16, wherein the protein comprises an antibody binding site.
 - 18. The method of claim 14, wherein the compound is a nucleic acid or mimetic.

- 19. The method of claim 18, wherein the nucleic acid or mimetic is an antisense, a ribozyme, an aptamer, or an interfering RNA.
- 5 20. The method of claim 14, wherein the compound is an organic molecule less than 1000 Dalton.
 - 21. The method of claim 20, wherein the compound comprises the following structure I or II

$$R_2$$
 N
 X
 R_1

$$R_2$$
 R_3
 R_1
 R_1
 R_1

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- 22. A kit comprising
- (a) a pharmaceutical composition comprising the agent used to inhibit MIF in claim 1, and
- (b) instructions for administering the composition to the mammal, wherein the mammal has type 1 diabetes or is at risk for type 1 diabetes.
 - 23. A kit comprising
- (a) a pharmaceutical composition comprising the agent used to inhibit MIF in claim 12, and
- (b) instructions for administering the composition to the mammal,wherein the mammal has type 1 diabetes or is at risk for type 1 diabetes.
- 24. Use of the agent used to inhibit MIF in claim 1 for the manufacture of a medicament for the treatment of a mammal having type 1 diabetes or at risk for type 1
 diabetes.
 - 25. Use of the agent used to inhibit MIF in claim 12 for the manufacture of a medicament for the treatment of a mammal having type 1 diabetes or at risk for type 1 diabetes.